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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
10/789,817	02/27/2004	John G. Babish	068911-0074	5656
23630 7590 12/23/2008 MCDERMOTT WILL & EMERY LLP 28 STATE STREET BOSTON, MA 02109-1775				
EXAMINER CARTER, KINDRA D				
ART UNIT 1617		PAPER NUMBER		
MAIL DATE 12/23/2008		DELIVERY MODE PAPER		

**Please find below and/or attached an Office communication concerning this application or proceeding.**

The time period for reply, if any, is set in the attached communication.

### Office Action Summary

**Application No.**

10/789,817

**Applicant(s)**

BABISH ET AL.

**Examiner**

KENDRA D. CARTER

**Art Unit**

1617

-- The MAILING DATE of this communication appears on the cover sheet with the correspondence address --  
**Period for Reply**

A SHORTENED STATUTORY PERIOD FOR REPLY IS SET TO EXPIRE 3 MONTH(S) OR THIRTY (30) DAYS, WHICHEVER IS LONGER, FROM THE MAILING DATE OF THIS COMMUNICATION.

- Extensions of time may be available under the provisions of 37 CFR 1.136(a). In no event, however, may a reply be timely filed after SIX (6) MONTHS from the mailing date of this communication.
- If NO period for reply is specified above, the maximum statutory period will apply and will expire SIX (6) MONTHS from the mailing date of this communication.
- Failure to reply within the set or extended period for reply will, by statute, cause the application to become ABANDONED (35 U.S.C. § 133). Any reply received by the Office later than three months after the mailing date of this communication, even if timely filed, may reduce any earned patent term adjustment. See 37 CFR 1.704(b).

**Status**

- 1) ☒ Responsive to communication(s) filed on 25 September 2008.
- 2a) ☒ This action is **FINAL**. 2b) ☐ This action is non-final.
- 3) ☐ Since this application is in condition for allowance except for formal matters, prosecution as to the merits is closed in accordance with the practice under *Ex parte Quayle*, 1935 C.D. 11, 453 O.G. 213.

**Disposition of Claims**

- 4) ☒ Claim(s) 1-32 is/are pending in the application.
- 4a) Of the above claim(s) 16-32 is/are withdrawn from consideration.
- 5) ☐ Claim(s) \_\_\_\_\_ is/are allowed.
- 6) ☒ Claim(s) 1-15 is/are rejected.
- 7) ☐ Claim(s) \_\_\_\_\_ is/are objected to.
- 8) ☐ Claim(s) \_\_\_\_\_ are subject to restriction and/or election requirement.

**Application Papers**

- 9) ☐ The specification is objected to by the Examiner.
- 10) ☐ The drawing(s) filed on \_\_\_\_\_ is/are: a) ☐ accepted or b) ☐ objected to by the Examiner.  
Applicant may not request that any objection to the drawing(s) be held in abeyance. See 37 CFR 1.85(a).  
Replacement drawing sheet(s) including the correction is required if the drawing(s) is objected to. See 37 CFR 1.121(d).
- 11) ☐ The oath or declaration is objected to by the Examiner. Note the attached Office Action or form PTO-152.

**Priority under 35 U.S.C. § 119**

- 12) ☐ Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f).
- a) ☐ All b) ☐ Some \* c) ☐ None of:
1. ☐ Certified copies of the priority documents have been received.
  2. ☐ Certified copies of the priority documents have been received in Application No. \_\_\_\_\_.
  3. ☐ Copies of the certified copies of the priority documents have been received in this National Stage application from the International Bureau (PCT Rule 17.2(a)).

\* See the attached detailed Office action for a list of the certified copies not received.

**Attachment(s)**

- 1) ☐ Notice of References Cited (PTO-892)
- 2) ☐ Notice of Draftsperson's Patent Drawing Review (PTO-948)
- 3) ☐ Information Disclosure Statement(s) (PTO/CDC)
- 4) ☐ Interview Summary (PTO-413)
- 5) ☐ Notice of Informal Patent Application
- 6) ☐ Other: \_\_\_\_\_
- Paper No(s)/Mail Date \_\_\_\_\_

### **DETAILED ACTION**

The Examiner acknowledges the applicant's remarks and arguments of September 25, 2008 made to the office action filed April 3, 2008. Claims 1-32 are pending. Claim 1 is amended and claims 16-32 are withdrawn.

For the reasons in the previous office action and below, the Applicant's arguments of the 35 USC 103(a) rejection of claims 1-15 were found not persuasive.

Due to the amendment to the claims, the modified 35 USC 103(a) rejection is made below. The Applicant's arguments are addressed below.

### ***Claim Rejections - 35 USC § 103***

The following is a quotation of 35 U.S.C. 103(a) which forms the basis for all obviousness rejections set forth in this Office action:

(a) A patent may not be obtained though the invention is not identically disclosed or described as set forth in section 102 of this title, if the differences between the subject matter sought to be patented and the prior art are such that the subject matter as a whole would have been obvious at the time the invention was made to a person having ordinary skill in the art to which said subject matter pertains. Patentability shall not be negated by the manner in which the invention was made.

Claims 1-15 are rejected under 35 U.S.C. 103(a) as being unpatentable over Ramirez (US 2002/0102345 A1) in view of Chappel et al. (Food and chemical Toxicology, 1998, vol. 36, pp. 915-922).

Ramirez teaches a beverage composition (i.e. oral and inherently possess a pharmaceutically acceptable carrier; addresses claims 14 and 15) comprising beer (see page 2, paragraph 29, line 3 and claim 1) and caffeine (see page 6, paragraphs 85-91 and claim 13; addresses claim 1, 7 and 9). Once the base composition is provided, special ingredients may then be incorporated into the liquid in such quantities and relative proportions that help enhance the body's alertness and energy sensation (see page 6, paragraph 82).

Ramierz does not teach tetra-hydroisalpha acid either derived or not derived from hops as disclosed in claims 1-6, nor the amounts of the compound nor ratio to the methylxanthine as disclosed in claims 8-13.

Chappel et al. teaches that during the storage of hops, deterioration may occur with losses in bittering value, thus tetrahydroisohumulones are manufactured from alpha acids, such as those elected by the applicant, that are more stable and provide more efficient bittering agent that can be added to beer late in the brewing process (see page 915, column 2, first full paragraph in its entirety).

To one of ordinary skill in the art at the time of the invention would have found it obvious and motivated to combine the composition of Ramierz and tetra-hydroisalpha acids as those disclosed in claims 1-6 because Chappel et al. teaches that these

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compounds are added to the brewing process of beer to provide a more stable and efficient bittering agent. Thus, in the brewing of the Ramierz beer, one skilled in the art would be motivated to add a more stable and efficient bittering agent in order to overcome the problem of the beer not having a true taste of the bitterness in beer.

To one of ordinary skill in the art at the time of the invention would have found it obvious and motivated to combine the composition of Ramierz in view of Chappel et al. and the amounts of the tetra-hydroisoalpha acid and methylxanthine as well as the ratio between the two compounds as disclosed in claims 8-13 because one skilled in the art would be able to adjust the amounts of these compounds in order to provide the desired effect of bitterness (tetra-hydroalpha acid) and energy boost (methylxanthine; caffeine). Ramiez teaches that once the base composition is provided, special ingredients may then be incorporated into the liquid in such quantities and relative proportions that help enhance the body's alertness and energy sensation (see page 6, paragraph 82).

In regards to the use of the composition claimed for the treatment of inflammation, the intended use does not get patentable weight. The reason for combining the prior art does not have to be the same as the Applicant. The claims are examined on the merits of a composition and not a method for treatment of inflammation.

In regards to methylxanthine and tetra-hydroisoalpha acids being in synergistic amounts, Ramierz in view of Chappel et al. teach a synergistic amounts because Chappel et al. teach that tetrahydroisohumulones are manufactured from alpha acids, such as those elected by the applicant, that are more stable and provide more efficient bittering agent that can be added to beer late in the brewing process (see page 915, column 2, first full paragraph in its entirety). Thus, by adding tetra-hydroisoalpha acids to the beer composition of Ramierz, a more efficient and stable bittering agent is provided.

### ***Response to Arguments***

Applicant's arguments filed September 25, 2008 have been fully considered but they are not persuasive.

The Applicant argues that Ramierez nor Chappel et al. do not teach or suggest the elements of "synergy" and "anti-inflammation". The combination of methylxanthine and reduced isoalpha acid has shown to be unpredictable in its effectiveness and its synergy.

The Examiner disagrees because the synergistic amounts are not commensurate to scope. The claims encompass any methylxanthine and any reduced isoalpha acid, tetrahydroisoalpha acid, or hexa-hydroisoalpha acid at any ratio. The specification

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demonstrates synergy with RIAA and curcumin in certain ratios (see page 44, table 5). For instance, RIAA alone has an IC50 of 0.81 but does not get better with the addition of curcumin. On the other hand curcumin alone has an IC50 of 1.4 and only gets better at a ratio of 3:2 (IC50 of 1.1) and 3:1 (IC50 1.3) of RIAA:Curcumin. Therefore, synergy does not exist with all of the compound combinations and ratios of the current claims. In regards to the anti-inflammatory effect of the composition, the Examiner notes that the intended use does not get patentable weight in composition claims. The claims are only treated on the merits as related to a composition.

### ***Conclusion***

Applicant's amendment necessitated the new ground(s) of rejection presented in this Office action. Accordingly, **THIS ACTION IS MADE FINAL**. See MPEP § 706.07(a). Applicant is reminded of the extension of time policy as set forth in 37 CFR 1.136(a).

A shortened statutory period for reply to this final action is set to expire THREE MONTHS from the mailing date of this action. In the event a first reply is filed within TWO MONTHS of the mailing date of this final action and the advisory action is not mailed until after the end of the THREE-MONTH shortened statutory period, then the shortened statutory period will expire on the date the advisory action is mailed, and any extension fee pursuant to 37 CFR 1.136(a) will be calculated from the mailing date of

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the advisory action. In no event, however, will the statutory period for reply expire later than SIX MONTHS from the date of this final action.

No claims allowed.

Any inquiry concerning this communication or earlier communications from the examiner should be directed to KENDRA D. CARTER whose telephone number is (571)272-9034. The examiner can normally be reached on 7:30 am - 4:00 pm.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Sreeni Padmanabhan can be reached on (571) 272-0629. The fax phone number for the organization where this application or proceeding is assigned is 571-273-8300.



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Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see <http://pair-direct.uspto.gov>. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free). If you would like assistance from a USPTO Customer Service Representative or access to the automated information system, call 800-786-9199 (IN USA OR CANADA) or 571-272-1000.

/K. D. C./  
Examiner, Art Unit 1617

/SREENI PADMANABHAN/  
Supervisory Patent Examiner, Art Unit 1617